IN THE CLAIMS:

11. (twice amended) A kit for detecting the presence or absence of one or more of a plurality nucleotide sequences in a sample, the kit comprising:

a capture agent; and

a plurality of electrophoretic probes selected from the group defined by the formula:

(D, M)-N-T

wherein:

(D, M)-N is an e-tag reporter released from an electrophoretic probe upon digestion of the electrophoretic probe by a nuclease, the e-tag reporter having a charge;

D is a detection group;

M is a non-oligomeric compound mobility modifier consisting of from 1 to 500 atoms selected from the group consisting of carbon, hydrogen, oxygen, phosphorus, nitrogen, sulfur, and boron;

N is a nucleotide; and

T is an oligonucleotide specific for at least one of the plurality of nucleotide sequences, each T having a length in the range of from 12 to 60 nucleotides such that at least one nucleotide of T has a capture ligand attached;

and wherein each e-tag reporter of the plurality of electrophoretic probes has a distinct charge/mass ratio so that the e-tag reporters form distinct peaks in an electropherogram upon electrophoretic separation;

and wherein the capture ligand specifically binds to the capture agent that has a charge opposite to that of e-tag reporters so that to exclude undigested electrophoretic probes are excluded from the electropherogram.

- 12. (previously amended) The kit of claim 11 wherein said formula is D-M-N-T and wherein M is a non-oligomeric compound consisting of from 1 to 300 atoms selected from the group consisting of carbon, hydrogen, oxygen, phosphorus, nitrogen, sulfur, and boron.
- 13. (previously presented) The kit of claim 12 wherein D is a fluorophore, chromophore, or an electrochemical label.

- 14. (previouly presented) The kit of claim 13 wherein said plurality is in the range of from 5 to 100.
- 15. (previouly amended) The kit of claim 14 further including a nuclease.
- 16. The kit of claim 14 wherein said fluorescent label is a fluorescein.
- 17. (previouly amended) The kit of claim 14 wherein said capture ligand is biotin wherein said capture agent is avidin or streptavidin.
- 18. (previouly amended) The kit of claim 14 further including said capture agent attached to a solid support.
- 19. (amended) A kit for detecting the presence or absence of one or more of a plurality nucleotide sequences in a sample, the kit comprising:
 - a capture agent; and
 - a plurality of electrophoretic probes selected from the group defined by the formula:

(D, M)-N-T

wherein:

(D, M)-N is an e-tag reporter released from an electrophoretic probe of the set upon digestion of the electrophoretic probe by a nuclease;

D is a detection moiety;

M is a non-oligomeric compound mobility modifier having a molecular weight of between 35 and 1500 daltons;

N is a nucleotide; and

T is an oligonucleotide specific for at least one of the plurality of nucleotide sequences, each T having a length in the range of from 12 to 60 nucleotides such that at least one nucleotide of T has a capture ligand attached;

and wherein each e-tag reporter of the plurality of electrophoretic probes has a distinct charge/mass ratio so that e-tag reporters of different electrophoretic probes form distinct peaks in an electropherogram upon electrophoretic separation;

and wherein the capture ligand specifically binds to the capture agent to exclude undigested electrophoretic probes from the electropherogram.

- 20. (previously presented) The kit of claim 19 wherein D is a fluorophore, chromophore, or an electrochemical label.
- 21. (previously presented) The kit according to claim 20 wherein said formula is D-M-N-T and wherein said plurality is in the range of from 5 to 100.
- 22. (previously presented) The kit of claim 21 wherein said capture ligand is biotin and wherein said capture agent is avidin or streptavidin.
- 23. (previouly presented) The kit of claim 21 further including a nuclease.

24. (previously presented) The kit of claim 21 wherein said e-tag reporter is selected from the group consisting of the following compounds: